

WHAT IS CLAIMED:

1. A DNA construct comprising:
5 one or more operatively linked nucleic acid molecules,
wherein the nucleic acid molecule encodes a serine proteinase inhibitor isolated
from *Brassica oleracea* having antibiosis activity and
an operably linked to a heterologous DNA promoter and
an operably linked 3' regulatory region.
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2. A DNA construct according to claim 1, wherein the nucleic
acid molecule either: (a) has a nucleotide sequence of SEQ. ID. No. 1; (b) encodes
a protein having an amino acid sequence of SEQ. ID. No. 2; or (c) hybridizes to at
15 the DNA molecule having a nucleotide sequence of SEQ. ID. No. 1 under
stringent conditions characterized by a hybridization buffer comprising 1M NaCl,
50 mM Tris-HCl, pH 7.4, 10 mM EDTA, 0.1% sodium dodecyl sulfate, 0.2%
ficoll, 0.2% polyvinylpyrrolidone, 0.2% bovine serum albumin, 50 µm g/ml *E.*
coli DNA at a temperature of 56°C.
- 20 3. An expression system comprising:
the DNA construct according to claim 1.
4. A host cell transduced with the DNA construct according to
claim 1.
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5. A host cell according to claim 4, wherein the cell is selected
from the group consisting of a bacterial cell, a virus, a yeast cell, and a plant cell.
6. A host cell according to claim 5, wherein the cell is a plant
30 cell.
7. A host cell according to claim 5, wherein the cell is a
bacterial cell.

8. A transgenic plant transformed with a DNA construct according to claim 1.

9. A transgenic plant according to claim 8, wherein the nucleic acid molecule either: (a) has a nucleotide sequence of SEQ. ID. No. 1; (b) encodes a protein having an amino acid sequence of SEQ. ID. No. 2; or (c) hybridizes to at the DNA molecule having a nucleotide sequence of SEQ. ID. No. 1 under stringent conditions characterized by a hybridization buffer comprising 1M NaCl, 50 mM Tris-HCl, pH 7.4, 10 mM EDTA, 0.1% sodium dodecyl sulfate, 0.2% ficoll, 0.2% polyvinylpyrrolidone, 0.2% bovine serum albumin, 50 µm g/ml *E. coli* DNA at a temperature of 56°C.

10. A transgenic plant according to claim 8, wherein the plant is selected from the group consisting of Gramineae, Liliaceae, Iridaceae, Orchidaceae, Salicaceae, Ranunculaceae, Magnoliaceae, Cruciferae, Rosaceae, Leguminosae, Malvaceae, Umbelliferae, Labitatae, Solanaceae, Cucurbitaceae, Compositae, and Rubiaceae.

11. A transgenic plant seed transformed with a DNA construct according to claim 1.

12. A transgenic plant seed according to claim 11, wherein the nucleic acid molecule either: (a) has a nucleotide sequence of SEQ. ID. No. 1; (b) encodes a protein having an amino acid sequence of SEQ. ID. No. 2; or (c) hybridizes to at the DNA molecule having a nucleotide sequence of SEQ. ID. No. 1 under stringent conditions characterized by a hybridization buffer comprising 1M NaCl, 50 mM Tris-HCl, pH 7.4, 10 mM EDTA, 0.1% sodium dodecyl sulfate, 0.2% ficoll, 0.2% polyvinylpyrrolidone, 0.2% bovine serum albumin, 50 µm g/ml *E. coli* DNA at a temperature of 56°C.

13. A transgenic plant seed according to claim 11, wherein the plant is selected from the group consisting of Gramineae, Liliaceae, Iridaceae, Orchidaceae, Salicaceae, Ranunculaceae, Magnoliaceae, Cruciferae, Rosaceae,

Leguminosae, Malvaceae, Umbelliferae, Labitatae, Solanaceae, Cucurbitaceae, Compositae, and Rubiaceae.

14. A method of conferring resistance to insects to plants
5 comprising:
transforming a plant or plant seed with the DNA construct
according to claim 1 and
growing the transformed plant or plants produced from the
seeds of a transformed plant under conditions effective to impart resistance to
10 insects.
15. A method according to claim 14, wherein a transgenic plant
is provided.
16. A method according to claim 14, wherein a transgenic plant
seed is provided.
17. A method according to claim 14, wherein the serine
proteainase inhibitor either: (a) has a nucleotide sequence of SEQ. ID. No. 1; (b)
20 encodes a protein having an amino acid sequence of SEQ. ID. No. 2; or (c)
hybridizes to at the DNA molecule having a nucleotide sequence of SEQ. ID. No.
1 under stringent conditions characterized by a hybridization buffer comprising
1M NaCl, 50 mM Tris-HCl, pH 7.4, 10 mM EDTA, 0.1% sodium dodecyl sulfate,
0.2% ficoll, 0.2% polyvinylpyrrolidone, 0.2% bovine serum albumin, 50 µm g/ml
25 *E. coli* DNA at a temperature of 56°C.
18. A method according to claim 14, wherein the insects are
selected from a group consisting of the orders of Lepidoptera, Coleoptera,
Diptera, Homoptera, Hemiptera, Thysanoptera, and Orthoptera.
19. A method according to claim 14, wherein the insects are
Heliothis viresens (tobacco budworm) or *Heliocoverpa zea* (corn earworm).

20. A method according to claim 14, wherein the transgenic plant is selected from a group consisting of Gramineae, Liliaceae, Iridaceae, Orchidaceae, Salicaceae, Ranunculaceae, Magnoliaceae, Cruciferae, Rosaceae, Leguminosae, Malvaceae, Umbelliferae, Labitatae, Solanaceae, Cucurbitaceae,
5 Compositae, and Rubiaceae.

21. A method of conferring resistance to insects to plants comprising:
applying a serine proteinase inhibitor having antibiosis
10 activity to a plant or plant seed under conditions effective to confer resistance to insects.

22. A method according to claim 21, wherein a plant is provided.
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23 A method according to claim 21, wherein a plant seed is provided.

24. A method according to claim 21, wherein the serine
20 proteinase inhibitor is isolated from *Brassica oleracea*.

25. A method according to claim 24, wherein the serine
protease inhibitor has an amino acid sequence comprising SEQ. ID. No. 2.

25 26. A method according to claim 21, wherein the insects are selected from the group consisting of the Orders of Lepidoptera, Coleoptera, Diptera, Homoptera, Hemiptera, Thysanoptera, and Orthoptera.

27. A method according to claim 26, wherein the insects are
30 *Heliothis viresens* (tobacco budworm) or *Helicoverpa zea* (corn earworm).

28. A method according to claim 21, wherein the plant is selected from a group consisting of Gramineae, Liliaceae, Iridaceae, Orchidaceae,

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